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Examiner's suggestion to overcome this problem by applying the request for continuation, while patent is pending is highly appreciated.

Concerning hematopoietic stem cells (HSCs) I have presented the argumentation that the specification while enabling for maintenance and expansion of HSCs also refer to generation of HSCs. The argumentation was as follows: As shown in the specification, the sorted CD34 positive cells expressing CD38 marker represent a hematopoietic progenitor cell population, with round non-polar morphology. After incubation with ACA specific antibody, these cells turned completely into cells with spindle morphology giving rise to generation of asymmetrically dividing cells (CD34+/CD38-) e.g. hematopoietic stem cells. The generation of the asymmetrically dividing cell (CD34+/CD38-) with polar spindle morphology, starting from progenitor cells (CD34+/CD38-) with round non-polar morphology, is a result of action of ACA. Crosslinking of ACA with specific antibody to ACA, changes the previous distribution of rafts, leading finally to their accumulation at one side of the cell membrane (spindle) allowing for concentration of the receptors for interaction with the ligands. This reaction is crucial for the mechanism of action of ACA as well as for the self-renewal for stem cell and explained why I think that the method described in the specification refers to generation, expansion and maintenance,

Regards,

Zorica Becker-Kojic

Dr. Zorica Becker-Kojic

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SEP 07 2010

Patent Application No.: 10/579,462

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

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Examiner: **Thaian N. Ton**
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Application No.: 11/579,462

Leimen, September 2, 2010

Filed: November 3, 2006

For: **USE OF THE ACA GLYCOPROTEIN FOR OBTAINING / MAINTAINING PLURIPOTENT
NON-EMBRYONIC STEM CELLS**

Summary of record of interview held on 22 July 2010

Commissioner for Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Signature and Date

Dear Sir:

Zorica Becker-Kojic, Sept. 2, 2010

This is in response to the substance of telephonic interview held on 22 July 2010 between applicant Dr. Becker-Kojic and examiner Dr. Thaian N. Ton. Discussed are the outstanding rejections of record and Dr. Becker-Kojic's proposed response.

I brief, while the objections stated by Examiner concerning the activators other than using ACA-specific antibody, or GPI-anchored glycoprotein ACA are removed from the claims, and the argumentation regarding generation of pluripotent stem cells starting with hematopoietic progenitor cells (CD34+/CD38+) which could be probative for the argumentation that the method described in the specification enable for generation of pluripotent stem cells as well, cannot be presented to the Examiner because the work is under review in Nature and displaying these data would compromise the process of accepting it for publication, a request for continuation will be filed. The